



10/524050 #2
PCT/GB 2003 / 0 0 3 4 2 8

Rec'd PCT/PTO 08 FEB 2005

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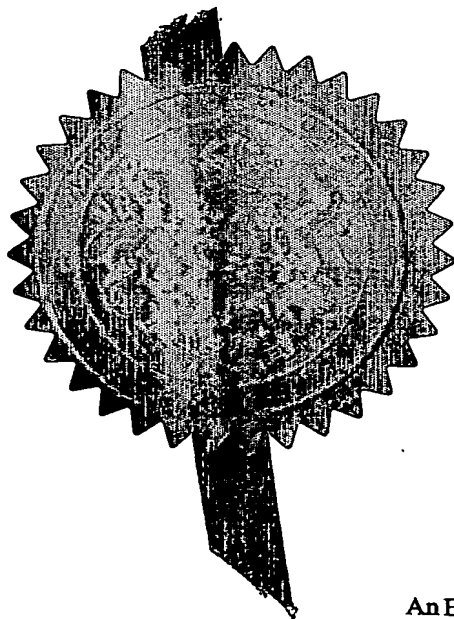
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16AUG02 E741448-1 C03022
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1. Your reference

9702

2. Patent application number

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0219037.9

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3. Full name, address and postcode of the or of each applicant (underline all surnames)

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Patents ADP number (if you know it)

If the applicant is a corporate body, give the country/state of its incorporation

6225916002

4. Title of the invention

PROCESS

5. Name of your agent (if you have one)

COLLINS, Frances Mary

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

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Country

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- a) any applicant named in part 3 is not an inventor, or
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11.

I/We request the grant of a patent on the basis of this application.

Signature F.M. Collins
COLLINS, Frances Mary

Date 15th August 200

12. Name and daytime telephone number of person to contact in the United Kingdom

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PROCESS

This present invention relates to oil field or gas field production chemicals, in particular, scale inhibitors and their use.

Scale inhibitors are used in production wells to stop scaling in the rock formation and/or in the production lines down hole and at the surface. Scale is a slightly soluble inorganic salt, such as barium or strontium sulphate, calcium carbonate, calcium sulphate or calcium fluoride. In the production of hydrocarbons from subterranean formations the deposition of scale on surfaces and production equipment is a major production problem. Scale build-up decreases permeability of the formation, reduces well productivity and shortens the lifetime of production equipment. In order to clean scaled-up wells and equipment it is necessary to stop the production i.e. by shutting in the well which is time-consuming and costly.

To minimise scale build-up a solution of a scale inhibitor may be injected by force into the formation via a production well-bore. After injection the production well is shut-in during which time the scale inhibitor is absorbed within the formation. After the shut-in period the production well is returned on stream and the inhibitor is slowly desorbed into the fluids in the formation to inhibit scale deposition. The fluids produced therefrom are analysed to determine the scale inhibitor concentration. When the concentration of inhibitor in the fluids has reduced to a certain level then further treatments will be required. An aqueous-based scale-inhibitor may have a short lifetime of a few weeks. The continual need for such treatments is therefore costly, not only in terms of production shut down periods but also in the cost of the chemical scale inhibitor used.

Other water-soluble or water-dispersible inhibitors used in production well environments include corrosion inhibitors, hydrogen sulphide scavengers or hydrate inhibitors. These too may need shut-ins.

According to US 5,547,025, it is well known to those skilled in the art that
5 gelled or crosslinked water-soluble polymers are useful in enhanced oil recovery and other oil field operations. In particular, they have been used to alter the permeability of underground formations in order to enhance the effectiveness of water flooding operations. Generally, polymers along with an appropriate crosslinking system are injected in an aqueous solution into the formation. The polymers then permeate into
10 and gel in the regions of the formation having the highest water permeability. Specifically, the process of US 5,547,025 comprises injecting into a formation a gelling composition which comprises a carboxylate-containing polymer, a crosslinking agent and a liquid wherein the gelling composition forms a gel when injected into the formation.

15 It has now been found that delayed release of an oil field or gas field production chemical can be achieved by incorporating the production chemical in a gelling composition which is injected into a hydrocarbon-bearing subterranean formation thereby allowing a decrease in the frequency of squeeze/shut in operations and an increase in the oil/gas production rate.

20 Thus, according to a first embodiment of the present invention there is provided a method of introducing an oil field or gas field production chemical into a subterranean formation comprising:
injecting a gelling composition comprising an aqueous liquid, an oil field or gas field production chemical, and a gellable polymer into said subterranean formation through a
25 well bore wherein the gellable polymer forms a gel in the formation and the oil or gas field production chemical is encapsulated in the gel; and
controllably releasing the oil or gas field production chemical into the formation fluids through breaking of the gel.

Controlled release of the oil field or gas field production chemical (hereinafter
30 "production chemical") upon breaking of the gel is advantageous in that it extends the lifetime of the production chemical and reduces the number of treatments required. Consequently production downtime and chemical costs are reduced.

It is envisaged that the gelling composition may comprise an admixture of the aqueous liquid, the production chemical and the gellable polymer and it is this admixture which is injected into the subterranean formation. Suitably, the production chemical may be dissolved, dispersed or suspended in the aqueous liquid of the admixture. Suitably, the gellable polymer is dissolved or dispersed in the aqueous liquid of the admixture.

Gellable polymers suitable for use in this invention are those which are soluble or dispersible in an aqueous liquid to increase the viscosity of the liquid. Preferred gellable polymers are those which are capable of crosslinking with a suitable crosslinking agent via crosslinkable groups.

Thus, according to a preferred embodiment of the present invention there is provided a method of treating a subterranean formation comprising: injecting a gelling composition comprising: (i) an aqueous liquid, (ii) an oil or gas field production chemical, (iii) a gellable polymer having crosslinkable groups, and (iv) a crosslinking agent into said subterranean formation through a well bore wherein a gel is formed in the formation through crosslinking of the crosslinkable groups of the polymer with the crosslinking agent and the oil or gas field production chemical is encapsulated in the gel; and controllably releasing the oil or gas field production chemical into the formation fluids through breaking of the gel.

It is envisaged that the gelling composition may comprise an admixture of the aqueous liquid, the production chemical, the gellable polymer and the crosslinking agent. Alternatively, the gellable polymer may be dispersed or dissolved in a first aqueous fluid (hereinafter "polymer slug") and the crosslinking agent may be dissolved in a second aqueous fluid (hereinafter "crosslinking agent slug"). The polymer slug and crosslinking agent slug are then injected sequentially into the formation in any order. Optionally, an aqueous spacer is injected between the polymer slug and crosslinking agent slug. Optionally, the subterranean formation is pre-flushed with an aqueous fluid prior to injection of the polymer slug and crosslinking slug. Suitably, the production chemical is dissolved, dispersed or suspended in one or more of the aqueous pre-flush fluid, the polymer slug, the crosslinking agent slug or aqueous spacer. During sequential addition, the gelling composition is formed within the formation by either back-producing the gellable polymer over the crosslinking agent or back-producing the crosslinking agent over

the gellable polymer. Where the production chemical is dissolved, dispersed or suspended in the pre-flush fluid it is necessary to back produce the pre-flush fluid over the polymer and crosslinking agent

Suitably the oil or gas field production chemical is selected from (i) scale inhibitors, (ii) corrosion inhibitors, (iii) hydrogen sulphide scavengers or (iv) hydrate inhibitors.

Scale inhibitors include water-soluble organic molecules having at least 2 carboxylic and/or phosphonic acid and/or sulphonic acid groups e.g. 2-30 such groups. Preferred scale inhibitors are oligomers or polymers, or may be monomers with at least one hydroxyl group and/or amino nitrogen atom, especially in hydroxycarboxylic acids or hydroxy or aminophosphonic, or, sulphonic acids. Scale inhibitors are used primarily for inhibiting calcium and/or barium scale. Examples of such compounds used as scale inhibitors are aliphatic phosphonic acids having 2-50 carbons, such as hydroxyethyl diphosphonic acid, and aminoalkyl phosphonic acids, e.g. polyaminomethylene phosphonates with 2-10 N atoms e.g. each bearing at least one methylene phosphonic acid group; examples of the latter are ethylenediamine tetra(methylene phosphonate), diethylenetriamine penta(methylene phosphonate) and the triamine- and tetramine-polymethylene phosphonates with 2-4 methylene groups between each N atom, at least 2 of the numbers of methylene groups in each phosphonate being different (e.g. as described further in published EP-A-479462, the disclosure of which is herein incorporated by reference). Other scale inhibitors are polycarboxylic acids such as acrylic, maleic, lactic or tartaric acids, and polymeric anionic compounds such as polyvinyl sulphonic acid and poly(meth)acrylic acids, optionally with at least some phosphonyl or phosphinyl groups as in phosphinyl polyacrylates. The scale inhibitors are suitably at least partly in the form of their alkali metal salts e.g. sodium salts.

Examples of corrosion inhibitors are compounds for inhibiting corrosion on steel, especially under anaerobic conditions, and may especially be film formers capable of being deposited as a film on a metal surface e.g. a steel surface such as a pipeline wall. Such compounds may be non-quaternised long aliphatic chain hydrocarbyl N-heterocyclic compounds; mono- or di-ethylenically unsaturated aliphatic groups e.g. of 8-24 carbons such as oleyl are preferred. The N-heterocyclic group can have 1-3 ring nitrogen atoms with 5-7 ring atoms in each ring; imidazole and imidazoline rings are

preferred. The ring may also have an aminoalkyl e.g. 2-aminoethyl or hydroxyalkyl e.g. 2-hydroxyethyl substituent. Oleyl imidazoline may be used. Where corrosion inhibitors are released into the formation using the method of the present invention, these inhibitors are effective in reducing corrosion of metal surfaces as they are produced out of the well.

Hydrogen sulphide scavengers include oxidants, such as inorganic peroxides, e.g. sodium peroxide, or chlorine dioxide, or aldehydes e.g. of 1-10 carbons such as formaldehyde or glutaraldehyde or (meth)acrolein.

Hydrate inhibitors include salts of the formula $[R^1(R^2)XR^3]^+Y^-$, wherein each of R^1 , R^2 and R^3 is bonded directly to X, each of R^1 and R^2 , which may be the same or different is an alkyl group of at least 4 carbons, X is S, NR^4 or PR^4 , wherein each of R^3 and R^4 , which may be the same or different, represents hydrogen or an organic group with the proviso that at least one of R^3 and R^4 is an organic group of at least 4 carbons and Y is an anion. These salts may be used in combination with a corrosion inhibitor and optionally a water soluble polymer of a polar ethylenically unsaturated compound. Preferably, the polymer is a homopolymer or a copolymer of an ethylenically unsaturated N-heterocyclic carbonyl compound, for example, a homopolymer or copolymer of N-vinyl-omega caprolactam. Such hydrate inhibitors are disclosed in EP 0770169 and WO 96/29501 which are herein incorporated by reference.

Suitably, the oil or gas field production chemical (hereinafter "production chemical") is water-soluble or water-dispersible. Alternatively, particles of the production chemical may be suspended in an aqueous liquid where the particles are sufficiently small that the particles disperse in the aqueous liquid component of the suspension. The particle size should also be sufficiently small to allow the particles to enter a formation. If the particles are too large they will tend to settle out and potentially lead to agglomeration problems. The particle size may be 100% less than 10 microns, preferably 100% less than 7 microns and especially 100% less than 5 microns. Preferably the particle size is not less than 25 nanometres and advantageously not less than 200 nanometres. The average particle size is usually between 1 and 3 microns. The particulate production chemical may be as described in EP 0902859 which is herein incorporated by reference.

The particulate production chemical may be coated with a suitable coating agent

also as described in EP 0902859. The coated particulate production chemical may be isolated from its production medium before dispersion in the gelling composition.

Suitable coating agents include water-soluble polymers or oil-soluble polymers.

Preferred water-soluble polymers for coating the particles of production chemical

5 include polyacrylic acids; polymaleic acids; polyacrylamide; polymethacrylate; polyvinylsulphonates; copolymers of monomers selected from the group consisting of acrylic acid, maleic acid, acrylamide, methacrylate, 2-acrylamido-2-methylpropane-sulfonic acid, and vinylsulphonate; lignosulphonates; hydroxy methyl cellulose; carboxy methyl cellulose; carboxy methyl ethyl cellulose; hydroxy methyl ethyl
10 cellulose; hydroxyl propyl methyl cellulose; methyl hydroxy propyl cellulose; sodium alginates; polyvinyl pyrrolidone; polyvinyl pyrrolidone acrylic acid co-polymers; polyvinyl pyrrolidone carolactam co-polymers; polyvinyl alcohol; polyphosphates, polystyrene-maleinates, poloxamers and poloxamines. Suitably, the poloxamers are linear ABA block co-polymers having the general structure $(EO)_n-(PO)_m-(EO)_n$ where n
15 and m are integers and EO and PO represents structural units derived from ethylene oxide and propylene oxide respectively. Suitably, the poloxamines are ABA block co-polymers having a branched structure with a central ethylene diamine bridge i.e. $[(EO)_n-(PO)_m]_2-N-CH_2-CH_2-N-[(EO)_n-(PO)_m]_2$ where n , m , EO and PO have the same meaning as for the poloxamers. Preferably, the water-soluble polymer has a molecular
20 weight in the range 1,000-100,000, preferably 5,000 to 30,000, for example, 15,000 to 25,000. Preferred oil-soluble polymers for coating the particles of production chemical include polyethers, polyamine derivatives or carbon backbone polymers having pendant nitrogen and/or oxygen atoms as described in EP 0902859.

Without wishing to be bound by any theory, the polymer will precipitate onto
25 the particles of production chemical and will at least partially coat the particles.

Suitably, at least 75%, preferably, at least 90%, more preferably, at least 95% of the surface of the particles is coated with the polymer. Preferably, the coating is continuous (100% surface coverage). Preferably, the coating has a thickness of less than 30 nm, preferably, less than 20 nm.

30 The coated production chemical provides for further controlled release of the production chemical into the formation fluids and/or the produced fluids.

Gellable polymers suitable for use in this invention include but are not limited to

biopolysaccharides, cellulose ethers and acrylamide-containing polymers. Suitably, the gellable polymers contain crosslinkable groups such as carboxylate, phosphonate or hydroxyl groups. Where the polymer contains carboxylate and/or phosphonate groups these groups may be in their acid or salt form. Suitable salts include ammonium, alkali metal or alkaline earth metal salts.

Suitable biopolysaccharides include natural and derivatized polysaccharides which are soluble or dispersible in an aqueous liquid to increase the viscosity of the liquid. Examples of natural gums include arabic gum, ghatti gum, tamarind gum, tagacanth gum, guar gum, locust bean gum, karaya gum, xanthan gum, galactomannan gum and the like. Preferred biopolysaccharides have molecular weights in the range 200,000 to 3,000,000. Modified gums such as carboxyalkyl derivatives, for example, carboxymethyl guar, and hydroxyalkyl derivatives, for example, hydroxyethyl guar, hydroxypropyl guar can also be employed. Doubly derivatized gums such as carboxymethylhydroxyethyl guar, carboxymethylhydroxypropyl guar can also be used.

Modified celluloses and derivatives thereof can also be employed in the practice of the present invention, for example, cellulose ethers, esters and the like. In general, any of the water-soluble cellulose ethers can be used. These cellulose ethers include, among others, the various carboxyalkyl cellulose ethers, such as carboxyethyl cellulose and carboxymethyl cellulose; mixed ethers such as carboxyalkylhydroxyalkyl cellulose ethers, e.g., carboxymethyl hydroxyethyl cellulose; hydroxyalkyl celluloses such as hydroxyethyl cellulose and hydroxypropyl cellulose; alkylhydroxyalkyl celluloses such as methylhydroxypropyl cellulose; alkyl celluloses such as methyl cellulose, ethyl cellulose and propyl cellulose; alkylcarboxyalkyl celluloses such as ethylcarboxymethyl cellulose; alkylalkyl celluloses such as methylethylcellulose; hydroxyalkylalkyl celluloses such as hydroxypropylmethyl cellulose; and the like.

Other suitable gellable polymers include the various polyacrylamides and related polymers which are partially hydrolysed and which are water-soluble, such as those disclosed in US 3,749,172 and EP 0 604 988 (herein incorporated by reference). Examples of suitable polymers include the homopolymers and copolymers of acrylamide and methacrylamide. Also, suitable are water-soluble copolymers resulting from the polymerisation of acrylamide and/or methacrylamide with another ethylenically unsaturated monomer copolymerisable therewith, wherein sufficient acrylamide and/or

methacrylamide is present in the monomer mixture to impart water-solubility to the resulting copolymer. The ethylenically unsaturated monomer which is copolymerisable with the acrylamide and/or methacrylamide may be selected from the group consisting of acrylic acid, methacrylic acid, vinyl sulfonic acid, vinylbenzylsulfonic acid, vinylbenzenesulfonic acid, vinyl acetate, vinylpyridine, styrene, acrylonitrile, methyl acrylonitrile, vinyl alkyl ether, vinyl chloride, maleic anhydride, N-vinyl-2-pyrrolidone, 2-acrylamido-2-methylpropanesulfonic acid, N-vinyl-2-pyrrolidone, N-vinyl formamide, and the like. Particularly preferred polymers include copolymers of N-vinyl-2-pyrrolidone and acrylamide; terpolymers of 2-acrylamido-2-methylpropanesulfonic acid, acrylamide and N-vinyl-2-pyrrolidone; and copolymers of sodium-2-acrylamido-2-methylpropanesulfonate and acrylamide. Other suitable gellable polymers contain crosslinkable phosphonate groups, for example, polymers which incorporate phosphonic acid monomers. Particularly preferred are copolymers of vinyl phosphonic acid monomers and acrylamide, copolymers of vinyl phosphonic acid monomers and methacrylamide or copolymers of vinyl phosphonic acid monomers, acrylamide and methacrylamide. These copolymers may also incorporate one or more further comonomers selected from the group consisting of acrylic acid, methacrylic acid, vinyl sulfonic acid, vinylbenzylsulfonic acid, vinylbenzenesulfonic acid, vinyl acetate, vinylpyridine, styrene, acrylonitrile, methyl acrylonitrile, vinyl alkyl ether, vinyl chloride, maleic anhydride, N-vinyl-2-pyrrolidone, 2-acrylamido-2-methylpropanesulfonic acid, N-vinyl-2-pyrrolidone, N-vinyl formamide, and the like.

Where the polymer is an acrylamide or related polymer, the mole percent of structural units containing carboxylate and/or phosphonate groups in the polymer is generally in the range of from 0.01 to 75 mole %. It is preferred that the mole percent of structural units containing carboxylate and/or phosphonate groups is in the range of 0.1 to 45, preferably 0.1 to 25, and most preferably 0.1 to 10 mole %.

Generally, the molecular weight of the acrylamide or related polymer is in the range of 10,000 to 50,000,000, preferably in the range 100,000 to 20,000,000, more preferably 200,000 to 15,000,000.

Further gellable polymers for use in the present invention include graft copolymers prepared by reacting hydrophilic polymers with certain allyl or vinyl monomers having a crosslinkable substituent. For example, graft copolymers of hydrophilic polymers and

vinyl phosphonate are disclosed in US 5,701,956 which is herein incorporated by reference. The hydrophilic polymer may be selected from polyacrylamides, polymethacrylamides, partially hydrolysed polyacrylamide, partially hydrolysed polymethacrylamide, copolymers containing acrylamide, copolymers containing methacrylamide, hydroxyalkylcelluloses, guar gum and derivatives thereof and the like. Graft copolymers of cellulose derivatives are described in US 4,982,793 and US 5,067,565 which are herein incorporated by reference. Preferably, the cellulose derivative is a hydroxyalkyl cellulose, in particular, hydroxyethyl cellulose. The preferred grafting monomers include glyceryl allyl ether, 2,3-dihydroxypropylmethacrylate, vinyl phosphonic acid, allyl glycidyl ether and glycidyl methacrylate.

The concentration of the gellable polymer in the gelling composition is generally in the range of about 0.01 to 0.5 weight percent, preferably about 0.05 to 0.4 weight percent, more preferably 0.05 to 0.35 weight percent, for example, 0.15 to 0.35 weight percent. A relatively low concentration of gellable polymer is advantageous since this mitigates the risk of a rigid blocking gel being formed in the formation.

Preferably, the gelling composition contains a buffering agent. Preferably, the buffering agent has a buffering capacity at a pH of up to about 5.5, preferably in a pH range of 4.5 to 5.5. A typical buffering agent is sodium acetate/acetic acid. Where the gelling composition contains a buffering agent, the concentration of buffering agent will be dependent on the type of buffering agent employed and the buffering capacity of the rock formation. Generally, it is preferred that the buffering agent is present at a concentration in the range 0.001 to 10% by weight, preferably 0.01 to 1% by weight (based on the weight of the gelling composition).

As discussed above, the aqueous gelling composition may include a cross-linking agent to further enhance the development of viscosity by cross-linking the polymer. The cross-linking agent may comprise any of the well known polyvalent metal compounds which are capable of creating a cross-linked structure with the particular polymer utilized. The presently preferred polyvalent metal compound is a metal compound selected from the group consisting of zirconium compounds, titanium compounds, aluminum compounds, iron compounds, chromium compounds, hafnium compounds, niobium compounds and antimony compounds, preferably zirconium and titanium compounds. Examples of suitable multivalent metallic compounds include, but are not limited to, ammonium

zirconium carbonate, sodium zirconium carbonate, potassium zirconium carbonate, ammonium zirconium fluoride, ammonium zirconium chloride, zirconium ammonium citrate, zirconium chloride, tetrakis(triethanolamine)zirconate, zirconium carbonate, zirconyl ammonium carbonate, zirconium lactate, zirconium acetylacetonate, zirconium diisopropylamine, zirconium 2-ethylhexanoate, zirconium acetate, zirconium neodecanoate, zirconium complex of hydroxyethyl glycine, zirconium malonate, zirconium propionate, zirconium tartrate, ammonium titanium carbonate, titanium acetylacetonate, titanium ethylacetoacetate, titanium triethanolamine, ammonium titanium lactate, titanium chloride, titanium carbonate, ammonium titanium chloride, titanium acetylacetonate, titanium triethanolamine, chromium citrate, aluminum acetate, potassium pyroantimonate, and combinations of any two or more thereof. These compounds are commercially available. Preferably, the crosslinking agent is a zirconium lactate of formula $[\text{CH}_3\text{CH}(\text{OH})\text{CO}_2]_n\text{X}_m\text{Zr}$ wherein X is an mono-valent anion, for example, a halide (chloride, bromide, iodide or fluoride) or hydroxide, m and n are integers such that $m+n = 4$ and n is 1 to 4, preferably 3 or 4.

The concentration of crosslinking agent in the gelling composition may vary over a broad range of from 0.001 to 0.5 weight percent based on the concentration of the polyvalent metal. The concentration of crosslinking agent in the gelling composition is preferably in the range of 0.01 to 0.25 weight percent, more preferably 0.025 to 0.2 weight percent, most preferably 0.025 to 0.15 weight percent based on the concentration of the polyvalent metal.

The aqueous fluid which is used to prepare the gelling composition may be pure water, tap water, seawater, aquifer water, a synthetic brine or a produced brine.

In a typical treatment, an aqueous pre-flush fluid (for example, a brine or fresh water) may first be injected (from a surface injection facility) into the formation followed by the gelling composition (injected either as an admixture or sequentially, as discussed above) and an optional overflush fluid. Preferably, the gelling composition is introduced into an aqueous zone of the subterranean formation. However, it is also envisaged that the gelling composition maybe introduced into a hydrocarbon zone of the subterranean formation. Without wishing to be bound by any theory it is believed that the gel which is formed within the formation acts as a relative permeability modifier. Thus, hydrocarbon formation fluids diffuse or percolate through the gel at a faster rate

than aqueous formation fluids.

The well may then be shut-in for a short period of time of up to 50 hours, preferably 2 to 24 hours, for example, 5 to 15 hours before the well is put back on production. After the well is put back in production, the production chemical is released
5 into the formation fluids and into the produced fluids upon degradation of the gel. The produced fluids may be analysed, for example, at the surface to monitor the concentration of production chemical to determine the need for further treatments.

The amount of production chemical used is preferably in the range from 1-25% w/w of the gelling composition, more preferably from 5-15% w/w, most preferably
10 from 6-10% w/w. Within these ranges the amount used would depend upon the nature of the chemical used and its intended purpose.

Where the components of the gelling composition are injected simultaneously, the gelling composition may contain a delaying agent in order to mitigate the risk of the gel being set up prematurely, for example, in the wellbore. A gelation delaying agent is
15 defined herein as a chemical or mixture of chemicals which delays the rate of gelation. A delaying agent useful for the retardation of the rate of gelation is generally a carboxylic acid or salts thereof. A commonly known delaying agent can also be an amine that has more than one functional group and contains one or more hydroxyls and that can chelate the polyvalent metal moiety of the polyvalent metal compound. It is
20 envisaged that the oil field or gas field production chemical, in particular, a scale inhibitor, may itself act as a gelation delaying agent.

Suitably, the extent of gelation of the polymer is such that the gel which is formed within the formation is strong enough to encapsulate an aqueous solution or dispersion of the production chemical and/or particles comprising the production
25 chemical within its structure.

The gelation rate is generally longer than 1 hour, preferably longer than 2 hours, more preferably longer than 3 hours, most preferably longer than 4 hours, for example, longer than 10 hours.

The gel which is set up in the formation must be capable of breaking so as to
30 release the solution or dispersion of the production chemical or the particles comprising the production chemical into the formation fluids.

The gel may degrade under the conditions encountered in the formation. Thus, the

gel may be thermally degradable or biodegradable. Alternatively, the gelling composition may contain an effective amount of a gel breaker so as to provide controlled breaking of the gel. The gel breaker may be selected from mild oxidizing agents such as ammonium persulfate, potassium dichromate, potassium permanganate, peroxides, alkali metal chlorites and alkali metal hypochlorites. Alternatively, the gel breaker may be a borate. It is also envisaged that the production chemical, for example, a scale inhibitor may act as a sequestration agent for the polyvalent ion of the crosslinking agent thereby acting as a gel breaker. Where the gellable polymer is a polysaccharide or a cellulose ether derivative, enzymes may be used as breakers. Suitable enzymes are alpha and beta amylases, amyloglucosidase, oligoglucosidase, invertase, maltase, cellulase and hemicellulase. Acids (for example, peracids) or chelants (for example, ethylenediaminetetracetic acid) may also be used to break the gel.

The amount of breaker employed is that required to reduce the viscosity of the gelled composition to a preselected lower viscosity or to achieve a complete break within a desired period of time. The optimum or effective amount of breaker employed in the present invention depends on factors such as the desired life of the gel, the particular gellable polymer and its concentration, the particular breaker and the formation temperature as well as other factors. Typically, however, from about 0.1 to about 10 wt% of breaker is employed in the gelling composition. Preferably, the amount of breaker employed is such to achieve a desired break in from about 12 to 500 hours. It is envisaged that the breaker may be encapsulated in the gelling composition. Alternatively, the breaker may be contained in the aqueous pre-flush fluid and is subsequently back produced through the gel.

In particular, the present invention provides a method for increasing the effectiveness of a water soluble or water dispersible oil or gas field production chemical by reducing the number of squeezing and shut-in operations needed to increase the production rate from an oil well or gas well, said method comprising:

A) injecting a gelling composition comprising an aqueous liquid, an oil or gas field production chemical and a gellable polymer into said subterranean formation through a production well bore wherein the polymer forms a gel in the formation and the oil or gas field production chemical is encapsulated in the gel;

B) after injecting the gelling composition, optionally over-flushing the oil or gas bearing rock formation with an aqueous fluid or an oil;

C) subsequently, shutting-in the well for a period of time; and

D) controllably releasing the oil or gas field production chemical into the formation

5 fluids though degradation of the gel.

The gel is preferably formed in the near well bore region of the well such that the production chemical is controllably released in the near well bore region. By "near well bore region" is meant a radial distance of less than 100 feet, preferably less than 50 feet, more preferably, less than 30 feet from the well bore.

10 The rate of degradation of the gel will be dependent upon, amongst others parameters, the formation temperature, the formation pressure, water cut of the produced fluids, formation permeability, flow rate and depth of placement. Suitably the gel starts to thermally degrade at a temperature in the range 50 to 150°C, preferably 50 to 100°C.

15 The invention will now be illustrated by means of the following examples.

Examples

Preparation of Gelling Composition

Xanthan gum (Barazan DTM ex Baroid), 2.5 g, was weighed into a plastic weighing boat. 497.5g of filtered seawater (0.45 µm mesh), collected off the Dorset
20 coast, was weighed out into a 1 litre beaker. A stirrer pellet (typically 30mm in length) was placed in the beaker which was then placed on a magnetic stirrer. The stirrer was switched on and the stirrer speed increased until a vortex was created in the seawater. The xanthan gum powder was then gradually introduced into the vortex. The resulting mixture was stirred until all of the powder had completely dissolved. 50 ml aliquots of
25 this solution were then syringed into 4 oz powder jars.

2.5ml of a 20% active solution of scale inhibitor (Calnox ML3263TM, ex Baker Petrolite) in seawater was added to each sample jar. Once the scale inhibitor had been introduced, the lids were replaced, and the jars were shaken for 30 seconds.

1 ml of a buffer solution was then pipetted into each jar. The jar lids were
30 replaced and the jars were again shaken for 30 seconds. The buffer formulation employed was prepared by mixing:

1) 20% wt/vol glacial acetic acid in deionised water, and

2) 50% wt/vol sodium acetate trihydrate in deionised water,
at a ratio of 1.262:1.86 vol/vol.

A 7% active solution of sodium zirconium lactate crosslinking agent was prepared by diluting a 50% active solution supplied by MEL chemicals with seawater.

5 1.43 ml of the crosslinking agent solution was added to each powder jar, and the jars shaken for 30 seconds.

The viscosity of the samples was measured using a Brookfield viscometer (LADV1 having RVT and HAT spindle assemblies) immediately after addition of the crosslinking agent solution. Measurements were also made one hour and 16 hours after
10 addition of the crosslinking agent solution with the samples maintained at a temperature of 25 °C during aging by means of a Julabo™ water bath. The results of the viscosity measurements are given in Table 1.

Table 1 – Viscosity Measurements

Time (mins)	Viscosity (cP)		
	30.6 s ⁻¹	61.2 s ⁻¹	183.6 s ⁻¹
0	52.5	38.4	17.0
60	160.1	101.9	40.3
960	190.9	117.8	42.1

Sandpack Tests

The performance of the gelling composition was assessed using a sand pack test. In this test, a 1 metre metal column of internal diameter 1.27 cm was packed with sand (deconsolidated Clashach sandstone, acid washed and sieved to a 20/40 mesh size)
20 which was then saturated with simulated formation water. By comparing the dry weight of the column with the wet weight, the liquid pore volume of the pack was determined. The pack was then placed in an oven and was connected to a pumping system that allowed fluids to be injected into the pack at a known volumetric flow rate with fluid effluent exiting the pack via a back pressure regulator. The back pressure regulator
25 allowed the pack to be pressurised and heated to above the boiling point of water at ambient temperature. The fluid effluent stream was fed into a sample fraction collector thereby allowing known volumes of samples to be collected for subsequent analysis.

Test Procedure

A sandpack of known liquid pore volume (of between 16.5 and 16.9 ml) was placed in the oven assembly and heated to a temperature of 120°C with the back pressure regulator set to a pressure of 80 barg. When the sandpack had reached the test temperature, 0.25 pore volumes (*ca.* 10 ml) of treatment fluid was pumped into the sandpack at a rate of 60 ml/hour. The treatment fluid was either a simulated seawater solution of Calnox ML3263™ (ex Baker Petrolite) having the same active content of scale inhibitor as the gelling composition prepared as described above (control experiment) or the gelling composition prepared as described above left to age for one hour following addition of the crosslinking agent solution (experiment according to the present invention). 0.25 pore volumes (*ca.* 10 ml) of simulated seawater was then pumped into the sandpack at a flow rate of 60 ml/hour. The sandpack was then closed off and maintained at a temperature of 120°C for 24 hours. After the shut in period, the sandpack was physically turned around in the oven assembly so that fluids could be injected into the sandpack in the reverse direction to simulate 'back' flow of a production well. Simulated seawater was then injected into the sandpack at a flow rate of 60 ml/hour for up to 8 hours and either 5 ml or 10 ml samples of the fluid effluent stream were collected for analysis.

The concentration of scale inhibitor (Calnox ML3263™) in the eluted fluid effluent stream was determined by titrating a known volume of barium chloride solution into the samples and measuring the turbidity developed after 3 minutes using a Hach turbidity meter. The turbidity reading was converted to a scale inhibitor concentration by reference to a previously determined calibrations curve. The results of these sandpack tests are given in Table 2 below.

The composition of the simulated seawater employed in the above test procedure is as follows:

ion	Concentration (ppm)
Na ⁺	11010
K ⁺	460
Mg ²⁺	1368
Ca ²⁺	428
Sr ²⁺	8
Cl ⁻	19700
SO ₄ ²⁻	2960
HCO ₃ ⁻	124

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Table 2 – Sandpack Tests

Control		Gelling Composition	
Volume eluted (ml)	ML3263 Concentration (ppm)	Volume eluted (ml)	ML3263 Concentration (ppm)
5	0.00	10	0.00
10	0.00	20	0.00
15	0.00	30	0.00
20	0.00	40	0.00
25	0.00	50	0.00
30	1.34	60	0.00
35	19.05	70	0.00
40	18.12	80	0.00
45	28.37	90	0.00
50	48.87	100	0.00
55	60.05	110	1.34
60	253.59	120	2.28
65	8798.92	130	7.87
70	19980.43	140	10.66
75	25571.19	150	28.37
80	28366.57	160	32.09
85	38097.28	170	35.82
90	45551.62	180	41.41
95	41824.45	190	346.77
100	26502.98	200	859.25
105	21844.02	210	3064.29
110	11853.80	220	7194.84
115	2091.22	230	16253.26
120	1998.04	240	18116.85
125	812.66	250	18116.85
130	346.77	260	17185.05
135	25.57	270	15321.47
140	19.98	280	10662.50
145	18.12	290	4399.46
150	13.46	300	1604.08
155	8.80	310	1387.07
160	5.07	320	160.41
165	0.41	330	40.48
		340	19.98

The data show that the length of time over which the scale inhibitor is eluted from the pack is increased by the use of the gelling composition of the present invention.

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